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<b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>  (use as many sheets as necessary)				<b>Complete if Known</b>	
				Application Number	10/806,295
				Filing Date	March 22, 2004
				First Named Inventor	LaColla, <i>et al.</i>
				Group Art Unit	Unassigned
Examiner Name	Unassigned				
Attorney Docket Number	06171.105033 IDX 1008 DIV				
Sheet	1	of	2		

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U.S. PATENT DOCUMENTS						
Examiner- Initials *	Cite No. <sup>1</sup>	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number	Kind Code (if known)			
LW	AA	4,866,084	A	Gunasekera <i>et al.</i>	09-12-1989	
	AB	5,124,327	A	Greenlee <i>et al.</i>	06-23-1992	
	AC	5,489,685	A	Houpis <i>et al.</i>	02-06-1996	
	AD	5,527,819	A	Williams <i>et al.</i>	06-18-1996	
	AE	5,830,894	A	Pevear <i>et al.</i>	11-03-1998	
	AF	5,852,011	A	Matsunaga <i>et al.</i>	12-22-1998	
	AG	5,929,114	A	Domagala <i>et al.</i>	07-27-1999	
	AH	5,935,982	A	Dykstra <i>et al.</i>	08-10-1999	
	AI	5,945,440	A	Kleinschroth <i>et al.</i>	08-31-1999	
	AJ	5,981,525	A	Farina <i>et al.</i>	11-09-1999	
LW	AK	6,025,390	A	Farina <i>et al.</i>	02-15-2000	

FOREIGN PATENT DOCUMENTS								
Examiner Initials *	Cite No. <sup>1</sup>	Foreign Patent Document			Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T <sup>6</sup>
		Office <sup>2</sup>	Number	Kind Code <sup>3</sup> (if known)				
LW	AL	EP	0,530,907	A1	Merck & Co.	03-10-1993		
I	AM	WO	94/19321	A1	Merck & Co.; Theoharides	09-01-1994		
I	AN	WO	02/083126	A1	Idenix; Univ. Degli Studi di Cagliari	10-24-2002		
LW	AO	WO	04/014364	A1	Idenix	02-19-2004		

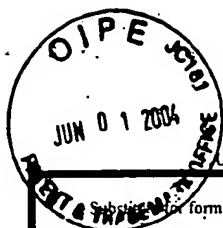
OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS			
Examiner Initials *	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>6</sup>
LW	AP	BALANI, S.K., <i>et al.</i> , "Biotransformation of 5-chloro-3-phenylthioindole-2-carboxamide (L-734,005) in rhesus monkeys and rat liver microsomes to a potent HIV-1 reverse transcriptase inhibitor," <i>Drug Metab. Dispos.</i> , 21(4):598-604 (July-August 1993).	

Examiner Signature	/Leonard Williams/ (06/26/2006)	Date Considered	
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\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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LW	AQ	CLAUSON-KAAS, N., <i>et al.</i> , "Preparation of <i>Cis</i> and <i>Trans</i> 2,5-Dimethoxy-2-(acetamidomethyl)-2,5-dihydrofuran, of <i>Cis</i> and <i>Trans</i> 2,5-Dimethoxy-2-(acetamidomethyl)-tetrahydrofuran and of 1-Phenyl-2-(acetamidomethyl)pyrrole," <i>Acta Chem. Scand.</i> , 6:667-670 (1952).	
	BA	ELMING, N., <i>et al.</i> , "The preparation of pyrroles from furans," <i>Acta Chem. Scand.</i> , 6:867-874 (1952).	
	BB	GAGLIARDI, S., <i>et al.</i> , "5-(5,6-Dichloro-2-indolyl)-2-methoxy-2,4-pentadienamides: novel and selective inhibitors of the vacuolar H <sup>+</sup> -ATPase of osteoclasts with bone antiresorptive activity," <i>J. Med. Chem.</i> , 41(10):1568-1573 (May 7, 1998).	
	BC	PAUWELS, R., <i>et al.</i> , "Potent and selective inhibition of HIV-1 replication in vitro by a novel series of TIBO derivatives," <i>Nature</i> , 343(6257):470-474 (February 1, 1990).	
	BD	PAUWELS, R., <i>et al.</i> , "Potent and highly selective human immunodeficiency virus type 1 (HIV-1) inhibition by a series of alpha-anilinophenylacetamide derivatives targeted at HIV-1 reverse transcriptase," <i>Proc. Natl. Acad. Sci. USA</i> , 90(5):1711-1715 (March 1, 1993).	
	BE	PHILLIPS, R.R., "The Japp-Klingemann Reaction," <i>Org. Reactions</i> , 10:143-178 (1959).	
	BF	ROMERO, D.L., <i>et al.</i> , "Bis(heteroaryl)piperazine (BHAP) reverse transcriptase inhibitors: structure-activity relationships of novel substituted indole analogues and the identification of 1-[(5-methanesulfonamido-1H-indol-2-yl)-carbonyl]-4-[3-[(1-methylethyl)amino]-pyridinyl]piperazine monomethanesulfonate (U-90152S), a second-generation clinical candidate," <i>J. Med. Chem.</i> , 36(10):1505-1508 (May 14, 1993).	
LW	BG	WILLIAMS, T.M., <i>et al.</i> , "5-chloro-3-(phenylsulfonyl)indole-2-carboxamide: a novel, non-nucleoside inhibitor of HIV-1 reverse transcriptase, <i>J. Med. Chem.</i> , 36(9):1291-1294 (April 30, 1993).	

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